



Sheet 1 of 5

<b>Form PTO-1449 Modified</b>		Docket No. <b>CELL-0086 (PA 446.3)</b>	Serial No. <b>09/450,999</b>
List of Patent and Publications Cited by Applicant (Use several sheets if necessary)		Applicant <b>John Robert Porter, et al.</b>	
U.S. Department of Commerce Patent and Trademark Office		Filing Date <b>November 29, 1999</b>	Group <b>1624</b>
<b>OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)</b>			
<i>Tu</i>	<b>AA</b>	Alhaique, F., et al., "Cyclisation of dinitriles by sodium alkoxides a new synthesis of naphthyridines," <i>Tetrahedron Letters</i> , <b>1975</b> , 3, 173-174	
<i>Tu</i>	<b>AB</b>	Ames, D.E., et al., "Condensation of $\beta$ -dicarbonyl compounds with halogenopyridinecarb-oxylic acids. A convenient synthesis of some naphthyridine derivatives," <i>J.C.S. Perkin I</i> , <b>1972</b> , 705-710	
<i>g1</i>	<b>AC</b>	Bodor, N., "Novel approaches in prodrug design," <i>Alfred Benzon Symposium</i> , <b>1982</b> , 17, 156-177	
<i>gr</i>	<b>AD</b>	Brooks, Peter C., et al., "Antiintegrin $\alpha\beta 3$ blocks human breast cancer growth and angiogenesis in human skin," <i>J. Clin. Invest.</i> , <b>1995</b> , 96, 1815-1822	
*	<b>AE</b>	Bundgaard, H., <i>Design of Prodrugs</i> , <b>1985</b> , Elsevier, Amsterdam	
*	<b>AF</b>	Katritzky, A.R., et al. (Eds.), <i>Comprehensive Organic Functional Group Transformations</i> , Pergamon, <b>1995</b>	
<i>Tu</i>	<b>AG</b>	Davies, S..G., et al., "Asymmetric synthesis of R- $\beta$ -amino butanoic acid and S- $\beta$ -tyrosine: homochiral lithium amide equivalents for Michael additions to $\alpha,\beta$ -unsaturated esters," <i>Tetra. Asymmetry</i> , <b>1991</b> , 2(3), 183-186	
<i>gr</i>	<b>AH</b>	Erle, D.J., et al., "Expression and function of the MadCAM-1 receptor, integrin $\alpha 4\beta 7$ , on human leukocytes," <i>J. Immunol.</i> , <b>1994</b> , 153, 517-528	
*	<b>AI</b>	<i>Encyclopedia of Reagents for Organic Synthesis</i> , John Wiley and Sons (eds.), <b>1995</b>	
<i>Tu</i>	<b>AJ</b>	Giacomello, et al., "Synthesis of 2,6-naphthyridine," <i>Tetra. Letters</i> , <b>1965</b> , 16, 1117-1121	
<b>EXAMINER</b> <i>Thomas Olliff</i>	<b>DATE CONSIDERED</b> <i>3/14/02</i>		

\* A copy of these references will not be forwarded to the U.S. Patent and Trademark Office since they are believed to be too voluminous and easily obtainable by the Examiner



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*	<b>AK</b>	Green, T.W., et al., "Protective Groups in Organic Synthesis," <i>John Wiley and Sons (eds.)</i> , 1991	
	<b>AL</b>	Hammes, H., et al., "Subcutaneous injection of a cyclic peptide antagonist of vitronectin receptor-type integrins inhibits retinal neovascularization," <i>Nature Medicine</i> , 1996, 2, 529-533	
	<b>AM</b>	Hodivala-Dilke, K.M., " $\beta$ 3-integrin-deficient mice are a model for glanzmann thrombasthenia showing placental defects and reduced survival," <i>J. Clin. Invest.</i> , 1999, 103(2), 229-238	
	<b>AN</b>	Kalvin, D.M., et al., "Synthesis of (4R)-D,L-[4- <sup>2</sup> H]- and (4S)-D,L-[4- <sup>2</sup> H] homoserine lactones," <i>J. Org. Chem.</i> , 1985, 50, 2259-2263	
	<b>AO</b>	Koivunen, E., et al., "Selection of peptides binding to the $\alpha_5\beta_1$ integrin from phage display library," <i>J. Biological Chemistry</i> , 1993, 268(27), 20205-20210	
	<b>AP</b>	Mitjans, F., et al., "An anti- $\alpha v$ -integrin antibody that blocks integrin function inhibits the development of a human melanoma in nude mice," <i>J. Cell Science</i> , 1995, 108, 2825-2838	
	<b>AQ</b>	Molina, P., et al., "Iminophosphorane-mediated annelation of a pyridine ring into a preformed pyridine one: synthesis of naphthyridine, pyrido [1,2-c] pyrimidine and pyrido [1,2-c] quinazoline derivatives," <i>Tetrahedron</i> , 1992, 48(22), 4601-4616	
	<b>AR</b>	Newham, P., et al., "Integrin adhesion receptors: structure, function and implications for biomedicine," <i>Molecular Medicine Today</i> , 1996, 304-313	
	<b>AS</b>	Numata, A., et al., "General synthetic method for naphthyridines and their N-oxides containing isoquinolinic nitrogen," <i>Synthesis</i> , 1999, 2, 306-311	
	<b>AT</b>	Sakamoto, T., et al., "Condensed heteroaromatic ring systems. III. synthesis of naphthyridine derivatives by cyclization of ethynylpyridinecarboxamides," <i>Chem. Pharm. Bull.</i> 1985, 33(3), 626-633	
<b>EXAMINER</b> <i>John Allard</i>		<b>DATE CONSIDERED</b> <i>3/14/02</i>	

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Sheet 3 of 5

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<b>OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)</b>				
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	AV	Srivatsa, S.S., et al., "Selective $\alpha\beta 3$ integrin blockade potently limits neointimal hyperplasia and lumen stenosis following deep coronary arterial stent injury: evidence for the functional importance of integrin $\alpha\beta 3$ and osteopontin expression during neointima formation," <i>Cardiovascular Research</i> , <b>1997</b> , 36, 408-428		
	AW	Stupack, D.G., et al., "induction of $\alpha_v\beta_3$ integrin-mediated attachment to extracellular matrix in $\beta_1$ integrin (CD29)-negative B cell lines," <i>Experi. Cell Research</i> , <b>1992</b> , 203, 443-448		
	AX	Tan R., et al., "Synthesis of 2, 6-naphthyridine and some of its derivatives," <i>Tetrahedron Letters</i> , <b>1965</b> , 31, 2737-2744		
	AY	Rico, J.G., et al., "A highly stereoselective michael addition to an $\alpha\beta$ -unsaturated ester as the crucial step in the synthesis of a novel $\beta$ -amino acid-containing fibrinogen receptor antagonist," <i>J. Org. Chem.</i> , <b>1993</b> , 58, 7948-7951		
	AZ	Zablocki, J.A., "Potent <i>in vitro</i> and <i>in vivo</i> inhibitors of platelet aggregation based upon the arg-gly-asp sequence of fibrinogen. (Aminobenzamidino)succinyl (ABAS) series of orally active fibrinogen receptor antagonists," <i>J. Med. Chem.</i> , <b>1995</b> , 38, 2378-2394		
EXAMINER	<i>Alvin Ottie Jr.</i>		DATE CONSIDERED	<i>3/14/01</i>



Sheet 4 of 5

Form PTO-1460 Modified		Docket No. <b>CELL-0086</b> <b>(PA 446.3)</b>	Serial No. <b>09/450,999</b>
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Examiner Initial		Document No.	Date	Country	Translation YES      NO
TEN	BA	WO 97/04247	02/06/97	PCT	
	BB	WO 97/23480	07/03/97	PCT	
	BC	WO 97/36858	10/09/97	PCT	
	BD	WO 97/36861	10/09/97	PCT	
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	BS	WO 99/52896	10/21/99	PCT	
	BT	WO 99/52898	10/21/99	PCT	
EXAMINER	<i>John Robert Porter</i>		DATE CONSIDERED	<i>3/14/02</i>	



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